

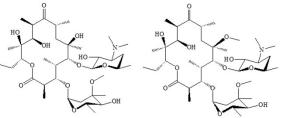


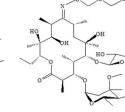
Supplementary information

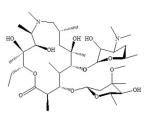
Preparation and application of molecularly imprinted 3 monolithic extraction column the selective for 4 microextraction of multiple macrolide antibiotics 5 from animal muscles 6

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Erythromycin

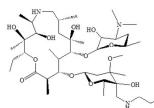
Clarithromycin

Florfenicol

Roxithromycin

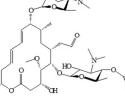
Azithromycin

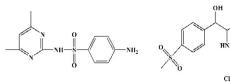
Tilmicosin

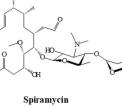


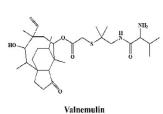
Tulathromycin

Sulfadimidine









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- Figure S1. Chemical structures of macrolide antibiotics, florfenicol, sulfadimidine and valnemulin

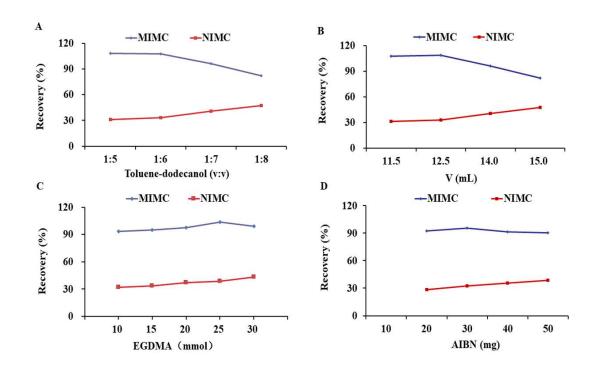
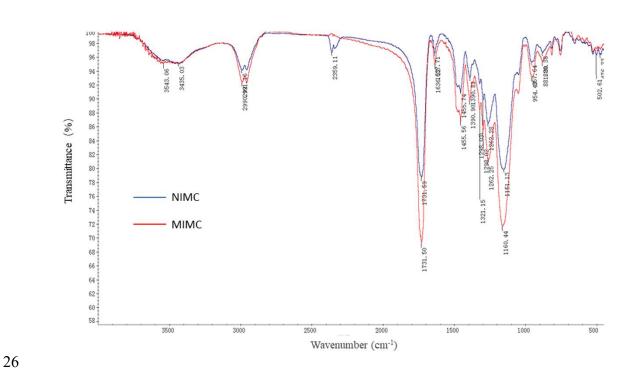
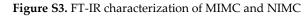




Figure S2. Effects of (A) different ratio of toluene to dodecanol as porogen, (B) porogen volume, (C) different amounts of EGDMA as cross-linker and (D) AIBN as initiator on the recovery of roxithromycin obtained from MIMC and NIMC





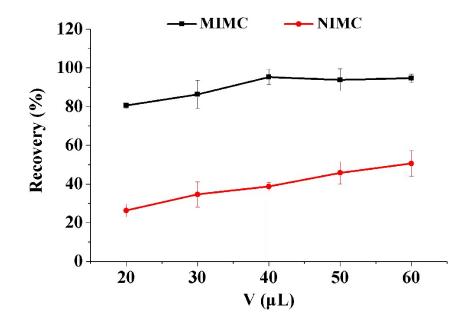


Figure S4. Effect of polymerization volume on the recovery of roxithromycin for MIMC and NIMC

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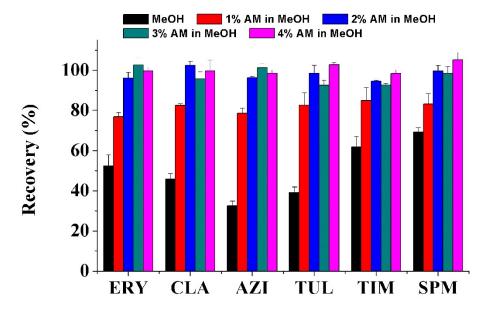
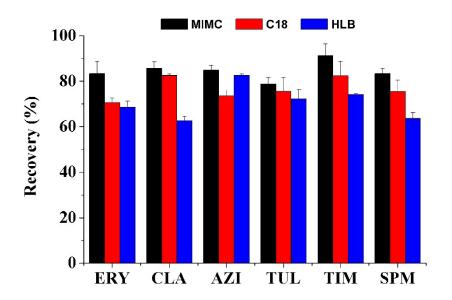


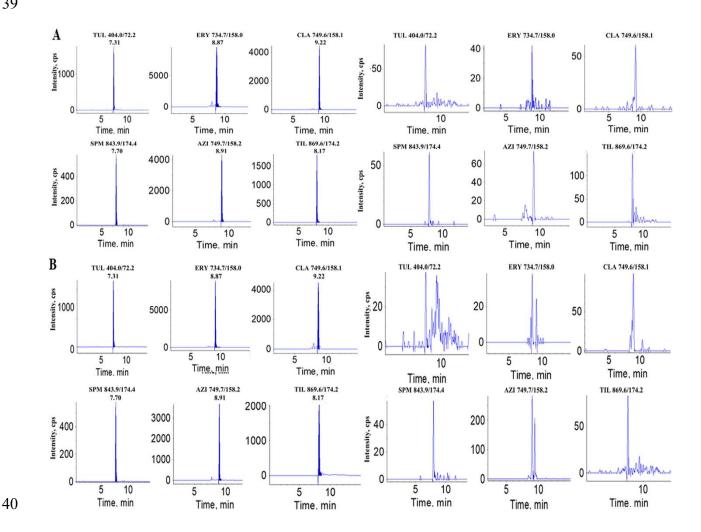


Figure S5. Effect of MeOH and different percentages of ammonium hydroxide (AM) in MeOH as elution
 solutions on the recoveries of macrolides drugs.



37 Figure S6. The comparison of MIMC, C18 and Oasis HLB cartridges on the recoveries of target macrolides (the 38 abbreviations are same as Figure 4) at 10 ng/mL spiked concentration of six macrolides in pork matrix

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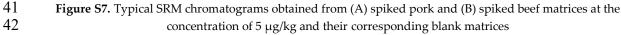


Table S1. SRM parameters for target analytes in positive ion mode a

Compounds	Abbr. ^b	Precursor ion	Product ion ^c	DP ^d	CE ^e	\mathbf{RT}^{f}
		[M + H] ⁺		(V)	(eV)	(min)
Tilmicosin	TIM	869.6	696.4*	130	60	8.26
			174.2	130	66	
Spiramycin	SPM	843.9	141.9*	110	50	7.85
			174.4	110	48	
Azithromycin	AZI	749.7	591.8*	80	46	9.11
			158.2	75	28	
Clarithromycin	CLA	749.6	591.5*	80	41	9.20
			158.1	80	26	
Erythromycin	ERY	734.7	576.5*	64	43	8.97
			158.0	64	27	
Tulathromycin	TUL	404	158.2*	71	33	7.37
			72.2	71	31	

^{*a*} SRM, selected reaction monitoring; ^{*b*} Abbr., abbreviations;

^c Product ion, the first product ion (*) of each analyte was used for quantification, and the second one was used for identification;

^d DP, declustering potential; ^e CE, collision energy; ^f RT, retention time.

Compounds	Matrix	Linear equation
Erythromycin	chicken	y=3.82×10 ³ χ+4.51×10 ²
	pork	$y=4.13 \times 10^{3} \chi + 1.62 \times 10^{2}$
	beef	$y=3.36\times10^{3}\chi+8.36\times10^{2}$
Clarithromycin	chicken	y=2.03×10 ³ χ+1.98×10 ³
	pork	y=3.85×10 ³ χ+4.23×10 ³
	beef	y=3.63×10 ³ χ+3.81×10 ³
Tulathromycin	chicken	$y=4.92 \times 10^{2} \chi+1.65 \times 10^{2}$
	pork	$y=5.19\times10^{2}\chi+4.26\times10^{2}$
	beef	$y=5.42 \times 10^{2} \chi + 3.67 \times 10^{2}$
Azithromycin	chicken	y=2.86×10 ³ χ+1.16×10 ²
	pork	$y=3.25\times10^{3}\chi+7.27\times10^{2}$
	beef	y=2.26×10 ³ χ+7.61×10 ²
Spiramycin	chicken	y=2.03×10 ² χ+1.81×10 ²
	pork	y=2.36×10 ² χ+2.53×10 ²
	beef	y=3.58×10 ² χ+1.32×10 ²
Tilmicosin	chicken	y=7.83×10 ² χ+3.35×10 ²
	pork	y=5.62×10 ² χ+4.12×10 ²
	beef	$y=6.15 \times 10^2 \chi + 4.69 \times 10^2$

Table S2. Linear equation for each analyte in three matrices